



MASSACHUSETTS GENERAL HOSPITAL
CENTER FOR INTEGRATED DIAGNOSTICS
DIAGNOSTIC MOLECULAR PATHOLOGY LABORATORY
55 FRUIT STREET, GRJ-1015
BOSTON, MA 02114

www.massgeneral.org/pathology/cid

MOLECULAR DIAGNOSTICS REQUISITION

- Label both the containers and this requisition with patient's name and ID
- Misabeled specimens will not be accepted

BILLING, SPECIMEN SUBMISSION, TESTING
STATUS QUESTIONS: 617-724-1285

TECHNICAL QUESTIONS: 617-643-2716

FAX: 617-643-1623

Date:	Location and Phone
Patient Identification are requirements: FULL NAME, MEDICAL RECORD NUMBER, SEX, DATE OF BIRTH	
Diagnosis/Differential Dx (required):	
PATHOLOGY LAB LABEL HERE	
Requesting the services below acknowledges an H&E review for sample adequacy. An interpretive report will be provided unless this box is checked <input type="checkbox"/>	

Date Collected:	Time Collected:	Completed by:
Requesting Physician Name (required):		MGH Provider # <input type="text"/>
Requesting Clinician Signature (required):		CLINICIAN: for tests indicated with an asterisk (*), please initial to attest that informed consent for testing has been obtained and documented in the patient's medical record. Requesting Clinician's Initials (required for *): <input type="text"/>
Path Resident (if applicable):		Pathology Staff:
Sample Origin (Institution, City, State, Phone)		Specimen Number/ID:
		Block ID/Slides:
BILLING: <input type="checkbox"/> ROUTINE <input type="checkbox"/> GLOBAL <input type="checkbox"/> OTHER		Private Consult Case: <input type="checkbox"/> YES <input type="checkbox"/> NO

ATTENTION: REQUIRED FOR ALL OUTPATIENTS - ALL APPLICABLE ICD-10 CODES (DX or SIGNS AND SYMPTOMS) FOR EACH TEST ORDERED. IF CODE(S) UNKNOWN, GO TO <http://www.icd10data.com>, OR PROVIDE TEXT ABOVE.

<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
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All test requests on surgical pathology specimens must include a surgical pathology report and diagnosis/indication. For outside specimens, please list outside specimen number/ID and origin.

TISSUE-BASED TESTING

For non-MGH requests include:

- FISH: H&E and 4 unstained 5 µm slides. Submit 2 additional unstained 5 µm slides for each additional FISH test.
- NGS/Non-NGS genotyping: H&E and 10 unstained 5 µm thick slides, unless otherwise noted.
- Consult Requisition Supplement for shipping information.

For MGH requests: the lab will obtain slides.

FISH

- | | |
|------------------------------------|---|
| <input type="checkbox"/> ALK | <input type="checkbox"/> HER2 (breast; WHO/ASCO 2018) |
| <input type="checkbox"/> ROS1 | <input type="checkbox"/> HER2 (non-breast) |
| <input type="checkbox"/> RET | <input type="checkbox"/> Ewing's Sarcoma (EWSR1) |
| <input type="checkbox"/> EGFR | <input type="checkbox"/> Myxoid Liposarcoma DDIT3 (CHOP) |
| <input type="checkbox"/> MET | <input type="checkbox"/> Synovial Sarcoma SS18 (SYT) |
| <input type="checkbox"/> FGFR1 | <input type="checkbox"/> Alveolar Rhabdomyosarcoma FOXO1 (FKHR) |
| <input type="checkbox"/> 1p/19q | <input type="checkbox"/> PDGFRA |
| <input type="checkbox"/> MYC/c-MYC | <input type="checkbox"/> KRAS |
| <input type="checkbox"/> BCL2 | <input type="checkbox"/> PIK3CA |
| <input type="checkbox"/> BCL6 | <input type="checkbox"/> CDKN2A |

Genotyping

NEXT GENERATION SEQUENCING (NGS) PANELS

NGS SOLID TUMOR

- ☐ NGS Solid Tumor Snapshot v2 (SNV/InDels/CNV); consent req*
- ☐ NGS Solid Fusion Assay v3 (ALK, ROS1, NTRK1/2/3, RET, MET, FGFR1/2/3, BRAF, NUTM1, MAML2, NRG1 and others) *

NGS Sarcoma Fusion Assay v1

NGS CELL FREE SNAPSHOT TESTING

- ☐ NGS cell-free Snapshot v1 (lung and breast) consent req*
- ☐ NGS cell-free Snapshot v1 (GI) consent req*

If submitting blood for cell free snapshot use 2 Streck tubes

HEMATOLOGY/LEUKEMIA

- ☐ HemeSnapshot v3 (SNV/InDels/CNV); consent req*
- ☐ HemeFusion assay v3

If submitting blood or bone marrow for SNaPshot, submit 3 mL

EDTA/purple top tube: **CORE:** place in **FRIDGE** for Molecular/J10

NON-NGS GENOTYPING

- ☐ Rapid BRAF (tissue: codon V600E/E2/D, V600K/R/M)
- ☐ Rapid IDH1/IDH2 (blood, tissue)
- ☐ Rapid EGFR Assay (tissue: MGH patients only)
- ☐ FLT3 (ITD)/ NPM1
- ☐ JAK2/CALR
- ☐ MLH1 Promoter Methylation
- ☐ MGMT Promoter Methylation
- ☐ Mismatch repair test: ☐ IHC(default) ☐ IHC+PCR ☐ PCR

Submit slides and H&E for BOTH tumor and normal tissues. When requesting PCR submit blood for normal control 3 mL EDTA/purple top tube: **CORE:** place in **FRIDGE** bin for Molecular/J10

BLOOD-BASED TESTING

TO BE DELIVERED TO CORE LAB, GRAY 5

☐ Array CGH* (documentation of consent required)

CORE: place in **FRIDGE** bin for Molecular/Jackson 10

- Submit 3 mL EDTA/purple top tube. Testing should only be ordered by a medical geneticist/genetic counselor. Please note that aCGH on prenatal samples is a send out test and will not be performed at MGH.

- Specify: ☐ Proband Family (specify relationship to the Proband in Notes section)
- Specify ICD-9 code below or in notes section:

- | | |
|--|---|
| <input type="checkbox"/> Multiple Congen. anomalies, NOS (759.7) | <input type="checkbox"/> CHD, unspecified (746.9) |
| <input type="checkbox"/> Hypotonia, congenital (779.89) | <input type="checkbox"/> Cleft palate, unspecified (749.00) |
| <input type="checkbox"/> Dysmorphic Features (744.89) | <input type="checkbox"/> Cleft lip, unspecified (749.10) |
| <input type="checkbox"/> Delayed Milestones (783.42) | <input type="checkbox"/> Skeletal anomalies, OS (756.0) |
| <input type="checkbox"/> Failure to Thrive (783.41) | <input type="checkbox"/> PDD - NOS (299.90) |
| <input type="checkbox"/> Macrocephaly (742.4) | <input type="checkbox"/> Autism (299.00) |
| <input type="checkbox"/> Microcephaly (742.1) | |

☐ Hemochromatosis* (documentation of consent required)

CORE: place in **FRIDGE** bin for Molecular/Jackson 10

- Submit 3 mL EDTA/purple top tube.

☐ Chimerism* (documentation of consent required)

CORE: place in **ROOM TEMP** bin for Molecular/Jackson 10

- Submit 2 ACD/yellow top tubes (PSoft Item ID #20303, BD tube ref #364606)

- ☐ Pre-transplant STR Genotyping
- ☐ Post-transplant Chimerism (requires pre-transplant genotyping)
Specify: ☐ Blood ☐ Bone Marrow

- Specify ICD-9 code below or in the ICD-9 section above:

- | | |
|--|---|
| <input type="checkbox"/> ALL, not having achieved remission, or failed (204.00) | <input type="checkbox"/> AML, not having achieved remission, or failed (205.00) |
| <input type="checkbox"/> Anemia, unspecified (285.9) | <input type="checkbox"/> Aplastic Anemia, other specified, other than constitutional (284.89) |
| <input type="checkbox"/> Aplastic Anemia (284.9) | <input type="checkbox"/> CLL, not having achieved remission, or failed (204.10) |
| <input type="checkbox"/> CML, not having achieved remission, or failed (205.10) | <input type="checkbox"/> Hodgkin's Lymphoma, site unspecified (201.90) |
| <input type="checkbox"/> Lymphoma - Non-Hodgkin's / B Cell / CNS, site unspecified (202.80) | <input type="checkbox"/> Lymphoma, Mantle cell, site unspecified (200.40) |
| <input type="checkbox"/> Multiple Myeloma, not having achieved remission, or failed (203.00) | <input type="checkbox"/> Myelodysplastic Syndrome, unspecified (238.75) |
| <input type="checkbox"/> Myelofibrosis, unspecified (289.83) | <input type="checkbox"/> T cell lymphoma, site unspecified (202.10) |

Other/Notes: